## AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

1-49. (Canceled).

50. (Currently Amended) A method for reducing restenosis following a vascular surgical procedure, the method comprising: locally administering to a human a biocompatible, nonbiodegradable sustained release dosage form comprising a cytostatic amount of a therapeutic agent comprising a cytostatic agent, an anti-migratory agent, a cytoskeletal inhibitor, or an anti-matrix agent dispersed in a polymer matrix, wherein said cytostatic amount of said eytostatie therapeutic agent, anti-migratory agent, cytoskeletal inhibitor, or anti-matrix agent inhibits a vascular smooth muscle cell activity without killing the cell, and wherein said eytostatic therapeutic agent, anti-migratory agent, eytoskeletal inhibitor, and anti-matrix agent are is not heparin, a radioisotope, a nitric oxide-releasing compound, suramin, methotrexate, adriamycin, a protein kinase inhibitor, staurosporin, an antisense oligonucleotide, colchicine, a peptidic inhibitor of a cellular factor that triggers proliferation of a smooth muscle cell or a pericyte, a growth factor inhibitor, a smooth muscle-derived growth factor inhibitor, an endothelial-derived growth factor inhibitor, a platelet homing receptor inhibitor, an integrin inhibitor, triazolopyrimidine, or a prostaglandin.

- 51. (Cancelled).
- 52. (Previously presented) The method of claim 50, wherein the vascular surgical procedure comprises placement of a stent.
- 53. (Previously presented) The method of claim 50, wherein the vascular surgical procedure comprises angioplasty.
- 54. (Previously Presented) The method of claim 50, wherein the locally administering comprises administering the cytostatic agent, anti-migratory agent, cytoskeletal inhibitor, or anti-matrix agent directly to vascular smooth muscle tissue.

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- 55. (Previously Presented) The method of claim 50, wherein the release of the cytostatic agent, anti-migratory agent, cytoskeletal inhibitor, or anti-matrix agent from the dosage form occurs during or after the vascular procedure.
- 56. (Previously Presented) The method of claim 50, wherein the cytostatic agent comprises a modified toxin, a radionuclide, a stimulator of the production or activation of TGF-beta, taxol, tamoxifen, TGF-beta, a nuclear enzyme DNA topoisomerase II inhibitor, a DNA polymerase inhibitor, an RNA polymerase inhibitor, an adenyl guanyl cyclase inhibitor, a superoxide dismutase inhibitor, a terminal deoxynucleotidyl-transferase, a reverse transcriptase, or lovastatin.
- 57. (Previously Presented) The method of claim 50, wherein the cytoskeletal inhibitor comprises a vinblastin, cytochalasin, taxol, taxotere, trichothecene, a modified diphtheria ricin toxin, or *Pseudomonas exotoxin*.
- 58. (Previously Presented) The method of claim 57, wherein the cytoskeletal inhibitor comprises taxol or taxotere.
- 59. (Currently Amended) The method of claim 57 50, wherein the sustained release dosage form is a microparticulate.
- 60. (Previously Presented) The method of claim 50, wherein the anti-migratory agent comprises a chemotactic factor inhibitor, a chemotactic factor receptor inhibitor, an intracellular cytoskeletal protein inhibitor, a caffeic acid derivative, nilvadipine, a steroid hormone, taxol, or cytochalasin.
- 61. (Previously Presented) The method of claim 50, wherein the anti-matrix agent comprises tamoxifen.
- 62. (Currently Amended) The method of claim 50, wherein the sustained release dosage form therapeutic agent comprises the cytostatic agent.
- 63. (Currently Amended) The method of claim 50, wherein the sustained release dosage form therapeutic agent comprises the cytoskeletal inhibitor.

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- 64. (Currently Amended) The method of claim 50, wherein the sustained release dosage form therapeutic agent comprises the anti-migratory agent.
- 65. (Currently Amended) The method of claim 50, wherein the sustained release dosage form therapeutic agent comprises the anti-matrix agent.